

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets

(11) Publication number:

0 334 244
A2

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 89104872.0

(51) Int. Cl.⁴: **C07K 7/18 , A61K 37/02 ,**
//C07K99/18

(22) Date of filing: 18.03.89

(30) Priority: 25.03.88 US 173391
01.03.89 US 315245

(43) Date of publication of application:
27.09.89 Bulletin 89/39

(84) Designated Contracting States:
AT BE CH DE FR GB IT LI LU NL SE

(71) Applicant: **THE PROCTER & GAMBLE**
COMPANY
One Procter & Gamble Plaza
Cincinnati Ohio 45202(US)

(72) Inventor: Gardner, Joseph Herman
4060 Boomer Road
Cincinnati, OH 45247(US)
Inventor: Correa, Paul Elliott
6511 Wesselman Road
Cincinnati, OH 45248(US)
Inventor: Berman, Elizabeth Faith
11787 Rose Lane
Cincinnati, OH 45246(US)
Inventor: Charest, Robert Paul
3935 Dust Commander Drive
Hamilton, OH 45011(US)

(74) Representative: Canonici, Jean-Jacques et al
Procter & Gamble European Technical
Center N.V. Temselaan 100
B-1820 Strombeek-Bever(BE)

(54) **Bradykinin Antagonist Peptides.**

(57) The present invention involves nona- or larger bradykinin antagonist peptides where certain amino acid residues of the nonapeptide hormone bradykinin or other substituted analogs of bradykinin are altered as follows: either position 2, position 3, or both consist of a L-acidic, L-amide or L-hydroxamate amino acid residue; position 7 consists of a D-aromatic amino acid residue; and position 4 preferably consists of a L-aliphatic or a D-cyclic amino acid residue.

EP 0 334 244 A2

5

BACKGROUND OF THE INVENTION

50 It is still another object of the present invention to provide novel methods for providing analgesia and/or smooth muscle relaxation, and for treating inflammation, rhinitis, rhinorrhea, sore throat pain, congestion and/or other conditions mediated by bradykinin, by administering such bradykinin antagonist peptides to humans or lower animals.

SUMMARY OF THE INVENTION

The present invention relates to bradykinin antagonist peptides having the structure:

5 H-L-Q-T-U-V-W-X-Y-Z-Arg-OH (1)

wherein

(a) L is selected from the group consisting of nil, a single basic amino acid residue of the D- or L-configuration, a peptide containing from about 2 to about 4 amino acid residues of the D- or L-configuration at least one of which is basic, and an N-terminal enzyme protecting group from the group consisting of
10 acyl-type protecting groups, aromatic urethane-type protecting groups, and alkyl-type protecting groups;

(b) Q is selected from the group consisting of Arg, Lys and Orn;

(c) T and U are each independently selected from the group consisting of D- and L-cyclic amino acid residues, D- and L-aliphatic amino acid residues, L-aromatic amino acid residues, L-acidic amino acid residues, L-amide amino acid residues, and L-hydroxamate amino acid residues; and wherein at least one
15 of T and U is selected from the group consisting of L-acidic amino acid residues and L-amide amino acid residues and L-hydroxamate amino acid residues;

(d) V is selected from the group consisting of D- or L-aliphatic, D- or L-cyclic, and D- or L-aromatic amino acid residues;

(e) W is selected from the group consisting of L-aliphatic and L-aromatic amino acid residues;

20 (f) X is selected from the group consisting of D- or L-aromatic and D- or L-aliphatic amino acid residues;

(g) Y is selected from the group consisting of D-aromatic amino acid residues; and

(h) Z is selected from the group consisting of L-aromatic and L-aliphatic amino acid residues; and the pharmaceutically acceptable salts thereof.

25

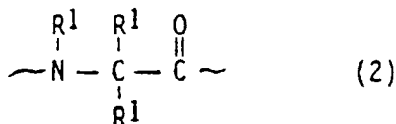
DETAILED DESCRIPTION OF THE INVENTION

30 The present invention involves peptides which are condensations of amino acid residues in particular sequences.

For purposes of this application, amino acid residues are categorized as aliphatic, cyclic, aromatic, basic, acidic, or amide based on the primary characteristic of their functional group.

As used herein, aliphatic amino acid residues have the following chemical structure:

35



40

wherein each R¹ is independently selected from the group consisting of hydrogen and saturated or unsaturated, straight or branched chain, unsubstituted or substituted alkyl having from 1 to about 6 carbon atoms. One R¹ attached to the carbon may be nil such that there is a double bond between the carbon and
45 the other R¹ attached to it. Substituted R¹ has substantially neutral (not acid or basic), non-aromatic substituents. Preferred R¹ are unsubstituted or substituted with hydroxy, alkoxy having from 1 to about 3 carbon atoms, halo or nitro; more preferred R¹ are unsubstituted or substituted with hydroxy. Preferred R¹ are C₁ to C₄ alkyl which are straight chain or have a single methyl branch. Preferred R¹ are saturated. Also preferred are aliphatic amino acid residues wherein the R¹ attached to the nitrogen is hydrogen and at least
50 one of the R¹'s attached to the carbon is a hydrogen.

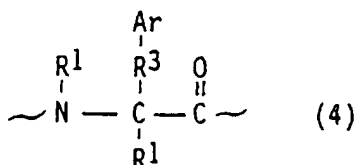
As used herein, cyclic amino acid residues have the chemical structure:

55



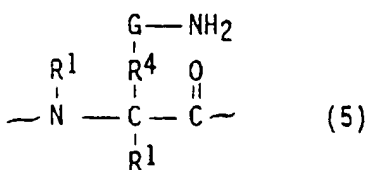
wherein R¹ is as defined hereinabove; wherein R² is saturated or unsaturated, straight or branched chain, unsubstituted or substituted alkyl having from about 1 to about 6 carbon atoms; wherein R⁷ is nil or CH₂; and wherein J is selected from the group consisting of nil, sulfur, and oxygen. J is preferably nil or sulfur; most preferably J is nil. R⁷ is preferably nil. R² is preferably C₂ to C₄, more preferably C₃ or C₄, most preferably C₃. R² is preferably saturated. R² is preferably straight chain. Substituted R² has substantially neutral, non-aromatic substituents. R² is preferably unsubstituted or substituted with hydroxy, alkoxy having from 1 to about 3 carbon atoms, halo or nitro; more preferably R² is unsubstituted or substituted with hydroxy; more preferably R² is unsubstituted. R¹ is preferably hydrogen, or when J is nil, R¹ may be nil such that there is a double bond between the carbon and R²; most preferably R¹ is hydrogen.

As used herein, aromatic amino acid residues have the chemical structure:



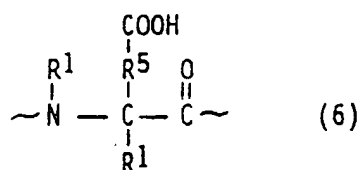
wherein each R¹ is as defined hereinabove; R³ is selected from the group consisting of nil, methylene and ethylene; and Ar is unsubstituted or substituted aryl. Each R¹ is preferably hydrogen. R³ is preferably methylene. Ar is preferably selected from the group consisting of unsubstituted and substituted phenyl, naphthyl, thienyl, pyridyl, indolyl and imidazolyl; most preferred Ar is unsubstituted or substituted phenyl. Substituted Ar has substantially neutral, non-aromatic substituents. Ar is preferably unsubstituted or substituted by hydroxy, C₁ to C₃ alkoxy, nitro or halo; more preferably Ar is unsubstituted or substituted with hydroxy; most preferably Ar is unsubstituted.

As used herein, basic amino acid residues have the chemical structure:

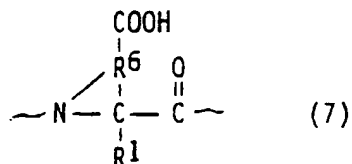


wherein each R¹ is as defined hereinabove; R⁴ is saturated or unsaturated, straight or branched chain, unsubstituted alkyl having from about 1 to about 6 carbon atoms; G is selected from the group consisting of nil and -NH-C(NH)-. Each R¹ is preferably hydrogen. R⁴ is preferably C₃ to C₄. R⁴ is preferably saturated. R⁴ is preferably straight chain.

As used herein, acidic amino acid residues have the chemical structure:



or



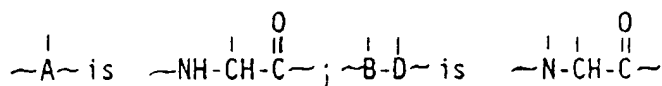
wherein each R^1 is as defined hereinabove; and R^5 and R^6 are saturated or unsaturated, branched or straight chain, unsubstituted alkyl having from 1 to about 6 carbon atoms. Each R^1 is preferably hydrogen. R^5 and R^6 are preferably C_1 to C_4 ; R^5 and R^6 are preferably saturated. R^5 and R^6 are preferably straight chain. R^5 is preferably C_1 to C_2 ; R^6 is preferably C_3 .

As used herein, amide amino acid residues have a chemical structure the same as acidic amino acid residues as defined hereinabove, but wherein the $-\text{COOH}$ is replaced by either $-\text{CONH}_2$ or $-\text{NHC(O)CH}_3$.

As used herein, hydroxamate amino acid residues have a chemical structure the same as amide amino acid residues as defined hereinabove having a $-\text{CONH}_2$ moiety, but wherein such $-\text{CONH}_2$ is replaced by $-\text{CONHOH}$.

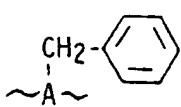
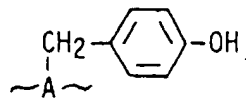
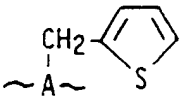
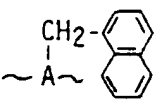
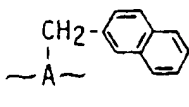
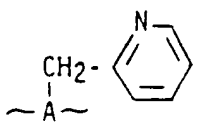
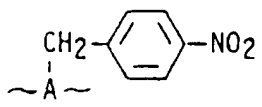
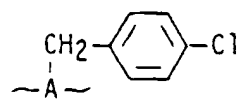
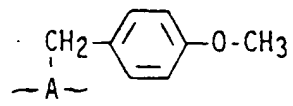
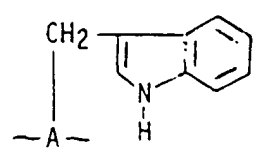
Table 1 hereinbelow provides examples of certain amino acid residues which are a part of the peptides of the present invention, abbreviations used in this application to denote such amino acid residues, and the chemical structures of these amino acid residues.

Table 1



Amino Acid Residue	Abbreviation*	Structure
<u>Aliphatic</u>		
glycyl	Gly	$ \begin{array}{c} \text{H} \\ \\ \sim \text{A} \sim \end{array} $
alanyl	Ala	$ \begin{array}{c} \text{CH}_3 \\ \\ \sim \text{A} \sim \end{array} $
valyl	Val	$ \begin{array}{c} \text{CH}(\text{CH}_3)_2 \\ \\ \sim \text{A} \sim \end{array} $
leucyl	Leu	$ \begin{array}{c} \text{CH}_2 - \text{CH}(\text{CH}_3)_2 \\ \\ \sim \text{A} \sim \end{array} $

Amino Acid Residue	Abbreviation*	Structure
<u>Aliphatic (continued)</u>		
isoleucyl	Ile	$\begin{array}{c} \text{H}_3\text{C}-\text{CH}-\text{C}_2\text{H}_5 \\ \\ \sim\text{A}\sim \end{array}$
seryl	Ser	$\begin{array}{c} \text{CH}_2\text{OH} \\ \\ \sim\text{A}\sim \end{array}$
threonyl	Thr	$\begin{array}{c} \text{CHOH}-\text{CH}_3 \\ \\ \sim\text{A}\sim \end{array}$
α -aminoisobutyryl	Aib	$\begin{array}{c} \text{CH}_3 \\ \\ \sim\text{NH}-\text{C}-\text{C}\sim \\ \quad \\ \text{CH}_3 \quad \text{O} \end{array}$
sarcosyl	Sar	$\begin{array}{c} \text{CH}_3 \quad \text{H} \\ \quad \\ \sim\text{B}-\text{D}\sim \end{array}$
<u>Cyclic</u>		
prolyl	Pro	$\begin{array}{c} \text{CH}_2-\text{CH}_2-\text{CH}_2 \\ \quad \quad \\ \sim\text{B} \quad \quad \text{D}\sim \end{array}$
hydroxyprolyl	Hyp	$\begin{array}{c} \text{CH}_2-\text{CHOH}-\text{CH}_2 \\ \quad \quad \\ \sim\text{B} \quad \quad \text{D}\sim \end{array}$
2-carboxy-thiazolidyl	Thz	$\begin{array}{c} \text{CH}_2-\text{CH}_2-\text{S} \\ \quad \quad \\ \sim\text{B} \quad \quad \text{D}\sim \end{array}$
2-carboxy-azetidyl	Azt	$\begin{array}{c} \text{CH}_2 \quad \quad \text{CH}_2 \\ \quad \quad \\ \sim\text{B} \quad \quad \text{D}\sim \end{array}$
pipecoliny1	Pip	$\begin{array}{c} \text{CH}_2-(\text{CH}_2)_2-\text{CH}_2 \\ \quad \quad \\ \sim\text{B} \quad \quad \text{D}\sim \end{array}$
2,3-dehydroprolyl	Epr	$\begin{array}{c} \text{CH}_2-\text{CH}_2-\text{CH} \quad \text{O} \\ \quad \quad \quad \\ \sim\text{N} \quad \quad \text{C} \quad \text{C}\sim \end{array}$

	<u>Amino Acid Residue</u>	<u>Abbreviation*</u>	<u>Structure</u>
5	<u>Aromatic</u>		
	phenylalanyl	Phe	
10	tyrosyl	Tyr	
15	β -(2-thienyl)alanyl	Thi	
20	β -(1-naphthyl)alanyl	1Nap**	
25	β -(2-naphthyl)alanyl	2Nap**	
30	β -(2-pyridyl)alanyl	Pyr	
35	p-nitrophenylalanyl	Nph	
40	p-chlorophenylalanyl	Cph	
45	O-methyltyrosyl	Mty	
50	tryptophyl	Trp	

55

Amino Acid Residue Abbreviation*

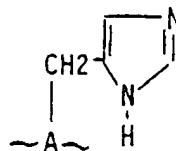
Structure

Aromatic (continued)

5

histidyl

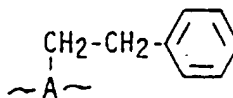
His



10

homophenylalanyl

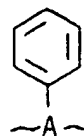
Hph



15

α -phenylglycyl

Pgl



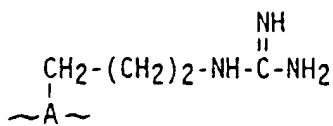
20

Basic

25

arginyl

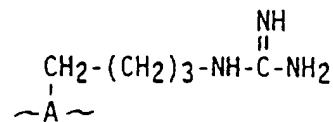
Arg



30

homoarginyl

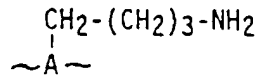
Har



35

lysyl

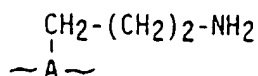
Lys



40

ornithyl

Orn

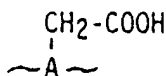


Acidic

45

aspartyl

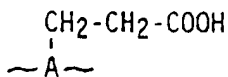
Asp



50

glutamyl

Glu



55

<u>Amino Acid Residue</u>	<u>Abbreviation*</u>	<u>Structure</u>
<u>Acidic (continued)</u>		
5 4-carboxypropyl	Cpr	$ \begin{array}{c} \text{COOH} \\ \\ \text{CH}_2 - \text{CH} - \text{CH}_2 \\ \quad \quad \\ \sim \text{B} \quad \quad \text{D} \sim \end{array} $
<u>Amide</u>		
10 asparaginy1	Asn	$ \begin{array}{c} \text{CH}_2 - \text{CONH}_2 \\ \\ \sim \text{A} \sim \end{array} $
15 glutaminy1	Gln	$ \begin{array}{c} \text{CH}_2 - \text{CH}_2 - \text{CONH}_2 \\ \\ \sim \text{A} \sim \end{array} $
20 4-carbamoylpropyl	Cmp	$ \begin{array}{c} \text{CONH}_2 \\ \\ \text{CH}_2 - \text{CH} - \text{CH}_2 \\ \quad \quad \\ \sim \text{B} \quad \quad \text{D} \sim \end{array} $
25 N-acetyl-4-aminopropyl	Aap	$ \begin{array}{c} \text{O} \\ \\ \text{HN} - \text{C} - \text{CH}_3 \\ \\ \text{CH}_2 - \text{CH} - \text{CH}_2 \\ \quad \quad \\ \sim \text{B} \quad \quad \text{D} \sim \end{array} $
30 N-acetylornithyl	Aor	$ \begin{array}{c} \text{O} \\ \\ \text{CH}_2 - (\text{CH}_2)_2 - \text{NH} - \text{C} - \text{CH}_3 \\ \\ \sim \text{A} \sim \end{array} $
<u>Hydroxamate</u>		
40 N-hydroxyasparaginy1	Nha	$ \begin{array}{c} \text{CH}_2 - \text{CONHOH} \\ \\ \sim \text{A} \sim \end{array} $
45 N-hydroxyglutaminy1	Nhg	$ \begin{array}{c} \text{CH}_2 - \text{CH}_2 - \text{CONHOH} \\ \\ \sim \text{A} \sim \end{array} $

50

55

 *Any three-letter abbreviation as provided in this Table 1 refers
 to an amino acid residue of the L-configuration, (except for

glycyl). An amino acid residue of the D-configuration is denoted
by the same three-letter abbreviation preceded by the letter "D".

**Collectively Nap

The present invention involves bradykinin antagonist nona-(or larger) peptides having the structure:

H-L-Q-T-U-V-W-X-Y-Z-Arg-OH (1)

Position: 0 1 2 3 4 5 6 7 8 9

In Structure (1), L is selected from the group consisting of nil; a single basic amino acid residue of the D- or L-configuration; a peptide containing from 2 to about 4 amino acid residues of the D- or L-configuration, at least one of which is a basic amino acid residue; and an N-terminal enzyme protecting group from the group consisting of acyl-type protecting groups, aromatic urethane-type protecting groups, and alkyl-type protecting groups such as, for example, acetyl, t-butyloxycarbonyl (Boc), and benzyloxycarbonyl. Preferred examples of L include Arg, DArg, Lys, DLys, Lys-Lys, Met-Lys, Gly-Arg-Met-Lys, DLys-Lys, Orn, Har, DHar, 5-guanidinopentanoyl, 6-guanidinohexanoyl, 6-aminohexanoyl, and 5-aminopentanoyl. More preferred L is selected from the group consisting of DArg, Lys-Lys and nil; most preferred L is Lys-Lys and especially DArg.

In structure (1), Q is selected from the group consisting of Arg, Lys and Orn. Most preferred Q is Arg.

In Structure (1), T and U are each independently selected from the group consisting of D- and L-cyclic amino acid residues, D- and L-aliphatic amino acid residues, L-aromatic amino acid residues, L-acidic amino acid residues, L-amide amino acid residues, and L-hydroxamate amino acid residues. In addition, at least one of T and U is selected from the group consisting of L-acidic amino acid residues, L-amide amino acid residues and L-hydroxamate amino acid residues. Preferred examples of T and U include Pro, DPro, Hyp, Azt, Thz, Inp, Epr, Asp, Glu, Asn, Gln, Cpr, Cmp, Aap, Aor, Nha and Nhg. It is preferred that one of T or U (preferably T) is selected from the group consisting of D- and L-cyclic amino acid residues, D- and L-aliphatic amino acid residues and L-aromatic amino acid residues, more preferably from Pro, DPro, Hyp, Azt, Thz, Inp and Epr, more preferably still from Pro and Hyp, most preferably Pro; while the other of T or U (preferably U) is selected from the group consisting of L-acidic amino acid residues, L-amide amino acid residues and L-hydroxamate amino acid residues, more preferably from Asp, Glu, Asn, Gln, Cpr, Cmp, Aap, Aor, Nha and Nhg, more preferably still from Asn, Cmp, Aap, Gln, Aor, Nha and Nhg; most preferably from Asn and Gln.

In Structure (1), V is selected from the group consisting of aliphatic, cyclic and aromatic amino acid residues of the D- or L-configuration. Preferred V is selected from the group consisting of Gly, Ala, DPro, DHyp and Sar; more preferred V are Gly and DPro; most preferred V is Gly.

In Structure (1), W is selected from the group consisting of aliphatic and aromatic amino acid residues of the L-configuration, preferably L-aromatic amino acid residues. Preferred examples of W include Phe, Thi, Nap, Pyr, Nph, Cph, Mty and Tyr. More preferred W is selected from the group consisting of Phe and Tyr; most preferred W is Phe.

In Structure (1), X is selected from the group consisting of aromatic and aliphatic amino acid residues of the D- or L-configuration. Preferred examples of X include Ser, Thr, Ala, Gly, DPhe, DThi, DNap, DPyr and Dcph. More preferred X is selected from the group consisting of Ser, Thr, Ala and Gly; most preferred X is Ser.

In Structure (1), Y is selected from the group consisting of aromatic amino acid residues of the D-configuration. Preferred examples of Y include DPhe, DTyr, DThi, DNap, DPyr, DNph, DCph, DMty, DTrp, DHis, DHph and DPgl; more preferred Y is selected from the group consisting of DPhe, DTyr, DTrp and DHis; most preferred Y is DPhe.

In Structure (1), Z is selected from the group consisting of aromatic and aliphatic amino acid residues of the L-configuration. Preferred examples of Z include Phe, Tyr, Pro, Thi, Pyr, Nap, Mty, Nph and Cph. More preferred Z is selected from the group consisting of Phe and Tyr; most preferred Z is Phe.

The synthesis of the peptides of Structure (1), including derivatization, activation, and coupling of protected amino acid residues, and their purification, and the analytical methods for determining the identity and purity are included in the general body of knowledge of peptide chemistry, as described, for example, in Kisfaludy, L., "Repetitive Methods in Solution", Chapter 6 of The Peptides, Vol. 2, Academic Press (pub.), 1980, for solution phase synthesis, and in Stewart, J.M. & J.D. Young, Solid Phase Peptide Synthesis, Freeman (pub.), 1969, for synthesis by the solid-phase method of Merrifield. A chemist skilled in

the art of peptide synthesis can synthesize the peptides of Structure (1) by standard solution methods or by manual or automated solid-phase methods.

5

EXAMPLES OF PEPTIDE SYNTHESIS

The following non-limiting examples further describe the peptides of the present invention and methods of making them.

10

Example 1

15

Preparation of H-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Asn³, DPhe⁷-bradykinin)

One gram of Boc-Arg(Tos)-polystyrene resin (Bachem Inc., Torrance, California) (Tos-tosyl) is loaded into the reaction vessel of a Biosearch 9500 automated peptide synthesizer. The peptide synthesis program is initiated by subjecting the amino acid-resin (0.30 mmol/g) to the Boc removal sequence as described below.

Boc Removal Sequence:

The amino acid-resin (or peptide-resin) is washed five times with methylene chloride (20 mL, 15 seconds per wash). The Boc group is removed by two successive treatments with 45% trifluoroacetic acid, 2.5% anisole in methylene chloride (60 seconds per treatment). The deprotected resin is then washed 2 times with methylene chloride and 3 times with a 1:1 mixture of methylene chloride:dimethylformamide and entered into the neutralization sequence.

Neutralization Sequence:

35

The trifluoroacetic salt of the amino acid-resin is washed two times with methylene chloride followed by three treatments with 10% diisopropylethylamine in methylene chloride. The amino acid resin containing the free amino group is washed 5 times with methylene chloride and then is ready for the coupling sequence.

40

Coupling Sequence:

The amino acid resin is washed once with a 2:1 mixture of methylene chloride:dimethylformamide. The coupling reaction is initiated with the addition of a 1:1 mixture of 0.6 M diisopropylcarbodiimide (Aldrich Chemical Company, Milwaukee, Wisconsin): 0.6 M Boc-L-phenylalanine (Bachem, Inc.) in methylene chloride (in 6.67 mole excess). The coupling reaction is mixed with bubbling nitrogen for 1 hour. After the reagents are removed by filtration the resin is washed successively with 1:1 methylene chloride:dimethylformamide (2X), methylene chloride (2X), 10% diisopropylethylamine (1X), methylene chloride (5X), and 1:1 methylene chloride:dimethylformamide. The resin is then recoupled for 1 hour (6.67 mole excess) with a 1:1 mixture of Boc-L-phenylalanine:diisopropylcarbodiimide. After removal of excess reagents the dipeptide-resin is washed twice with a 1:1 mixture of methylene chloride:dimethylformamide.

The Boc-Phe-Arg(Tos)-resin is then treated with the Boc-removal sequence described above followed by the neutralization sequence described above. The free amine-dipeptide-resin is then coupled with Boc-D-phenylalanine according to the coupling sequence described above to form the Boc-DPhe-Phe-Arg(Tos)-resin (tripeptide-resin).

The remaining amino acids are added in the following order: Boc-Ser, Boc-Phe, Boc-Gly, Boc-Asn, Boc-Pro, Boc-Arg(Tos) using the same Boc removal, neutralization and coupling sequences described above with the following exceptions: 1.5 equivalents of 1-hydroxybenzotriazole (Aldrich Chemical Company) are

added during the Asn and Arg couplings to suppress unwanted side reactions known to occur when coupling these residues.

After the nonapeptide (nine amino acid peptide) is assembled on the resin as described above the peptide is cleaved from the polystyrene resin using the following HF (hydrofluoric acid) reaction.

HF Cleavage Reaction:

The completed peptide-resin (2.3 gm) is suspended in 40 mL of condensed hydrofluoric acid containing 2 mL of anisole at -78°C . The reaction is stirred for 1 hour at -78°C followed by 1 hour at 0°C . The liquid HF is removed at 0°C under reduced pressure with a slow bleed of anhydrous argon gas passing through the reaction vessel. The free peptide is extracted from the resin by washing the resin-peptide mixture with 2N acetic acid. The acetic acid washes are combined and concentrated under vacuum to yield the crude peptide product.

The peptide is purified by first passing the crude peptide through a G-10 sephadex column (Sigma Chemical Company, St. Louis, Missouri), eluting with 1N acetic acid. Appropriate fractions are pooled after analysis by thin layer chromatography and then lyophilized to yield a white amorphous solid. The peptide is further purified using reverse phase chromatography on a C_{18} silica column (2.14 X 30 cm) (Rainin Instruments, Inc., Woburn, Massachusetts), eluting with a 30 minute gradient of 10-40% solvent B mixed with solvent A (solvent A = 0.1% trifluoroacetic acid in water, solvent B = 0.1% trifluoroacetic acid in acetonitrile). The appropriate fractions are pooled and lyophilized to yield the purified peptide product.

The peptide is characterized by mass spectral analysis on a Finnigan model TSQ-46C triple quadrupole instrument using the fast atom bombardment ionization technique from a glycerol matrix. The observed molecular ion, $\text{M} + \text{H} = 1127$, agrees with the calculated molecular weight for the desired structure. The synthetic peptide is also sequenced using Edman degradation sequencing techniques on an automated Applied Biosystems #420 sequencer to confirm the structure of the synthetic peptide.

Example 2

Preparation of H-Arg-Pro-Gln-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Gln³, DPhe⁷-bradykinin)

The peptide is made using the same methods described in Example 1 with the replacement of Gln for Asn in the coupling reaction for position 3 in the peptide sequence. The final peptide is analyzed by mass spectral analysis as described in Example 1 giving the correct molecular ion, $\text{M} + \text{H} = 1141$.

Example 3

Preparation of H-Arg-Pro-Asn-DPro-Phe-Ser-DPhe-Phe-Arg-OH (Asn³, DPro⁴, DPhe⁷-bradykinin)

The peptide is made using the same methods described in Example 1 with the replacement of DPro for Gly in the coupling reaction for position 4 in the peptide sequence. The final peptide is characterized by mass spectral analysis as described in Example 1 giving the correct molecular ion, $\text{M} + \text{H} = 1167$.

Example 4

Preparation of H-Arg-Asn-Pro-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Asn², DPhe⁷-bradykinin)

The peptide is made using the same methods described in Example 1 with the replacement of Asn for Pro and Pro for Asn in the coupling reaction for positions 2 and 3, respectively, in the peptide sequence. The final peptide is characterized by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1127$.

Example 5

Preparation of H-Arg-Pro-Asp-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Asp³, DPhe⁷-bradykinin):

The peptide is made using the same methods described in Example 1 with the replacement of Asp for Asn in the coupling reaction for position 3 in the peptide sequence. The final peptide is characterized by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1128$.

Example 6

Preparation of H-Arg-Pro-Aap-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Aap³, DPhe⁷-bradykinin) and the preparation of the amino acid Aap

Prior to the automated synthesis of the nonapeptide, the precursor of the unnatural amino acid residue N-acetyl-4-aminoprolyl (Aap) is prepared using the following procedure.

N-Boc-4-hydroxyproline benzyl ester:

N-tert-butyloxycarbonyl-4-hydroxyproline (Boc-Hyp) (Bachem, Inc.), is converted to its benzyl ester as follows. Boc-Hyp (20 g, 86 mmol) is dissolved in dimethylformamide (500 mL); potassium carbonate (5 equivalents) is added. The mixture is stirred vigorously while adding benzyl bromide dropwise over a period of 15 minutes. The reaction is heated to 55 °C and stirred for 1.5 hours at that temperature. After cooling to room temperature the reaction mixture is diluted with 1.5 L of ethyl acetate and placed in a separatory funnel. The ethyl acetate layer is washed sequentially with water (2 X 1.2 L), 0.5 N hydrochloric acid (1 L), water (1.5 L), and saturated sodium chloride (1.5 L). The organic layer is dried over magnesium sulfate, filtered and evaporated to dryness. The resulting crude product is purified using chromatography on silica gel column with 70:30 hexane: acetone as eluant. The appropriate fractions are pooled after TLC analysis yielding 22.8 g of clear viscous liquid (82%).

N-Boc-4-azidoproline benzyl ester:

The N-Boc-4-hydroxyproline (10 g, 31.2 mmole) from above is dissolved in dimethylformamide (14 mL), and triethylamine (10.7 mL) is added. The solution is chilled to 0 °C and methanesulfonyl chloride (mesyl chloride 4.1 mL) is added dropwise over a period of 12 minutes. The reaction is stirred at 0 °C for 1 hour followed by 1 hour at room temperature. The intermediate 4-O-mesyl proline is not isolated but is converted directly to the azide with the addition of sodium azide (15.2 g) into the above reaction mixture followed by stirring for 20 hours at room temperature. After the first 20 hours, an additional 15.2 g of sodium azide is added and the reaction is stirred for an additional 72 hours at room temperature. The reaction mixture is transferred to a separatory funnel and diluted with 1 L of water. The aqueous mixture is extracted twice with ethyl acetate (2 X 1 L) and the ethyl acetate extracts are combined and washed with water (1 L), 0.5 N HCl (1 L), water (1 L), saturated sodium chloride (1 L). The ethyl acetate layer is dried over magnesium sulfate and evaporated to dryness. The resulting crude product is purified on silica gel chromatography eluting with 8:2 hexane:acetone. The appropriate fractions are pooled after TLC analysis yielding 3.4 g of product

(showing a strong band in IR at 2100 cm^{-1}).

N-Boc-4-aminoproline:

5

The N-Boc-4-azidoproline benzyl ester from above (3.4 g, 9.8 mmol) is dissolved in methanol and placed under an inert atmosphere. Palladium on carbon (380 mg, 10%) is added and the reaction vessel is placed in a Parr hydrogenation apparatus and pressurized with 40 psi of hydrogen gas. The reaction vessel is shaken vigorously for 3 hours and then removed from the Parr apparatus. The resulting mixture is filtered
10 through celite and the methanol is removed in vacuo. The resulting white solid product is used directly in the following acetylation reaction.

α -N-Boc-N-(acetyl)-4-aminoproline:

15

The N-Boc-4-aminoproline from above (1.4 g, 6.1 mmol) is dissolved in 9 mL of 2.5 N NaOH, and 9 mL of *p*-dioxane is added. The resulting mixture is chilled in an ice bath to about 10°C , and acetic anhydride (1.0 mL) is added dropwise over a period of 10 minutes. The reaction is stirred for 30 minutes at 0°C , followed by stirring for 1 hour at room temperature. The reaction mixture is diluted with water (200 mL) and
20 extracted twice with ethyl ether. The aqueous layer is acidified to pH 2-3 with the addition of solid potassium hydrogen sulfate, then extracted with ethyl acetate (2 X 200 mL). The ethyl acetate extracts are combined, washed with saturated sodium chloride, dried over magnesium sulfate, and evaporated to dryness. The final product is analyzed by ^1H - and ^{13}C -NMR, mass spec. $M^{+} = 272$, providing the correct structure.

25

The N-acetyl-4-aminoproline is incorporated into position 3 of the peptide sequence indicated in this example above using the same methods of peptide synthesis described in Example 1. The structure of the nonapeptide is confirmed by mass spectral analysis, $M+H = 1167$, and amino acid sequencing as described in Example 1.

30

Example 7

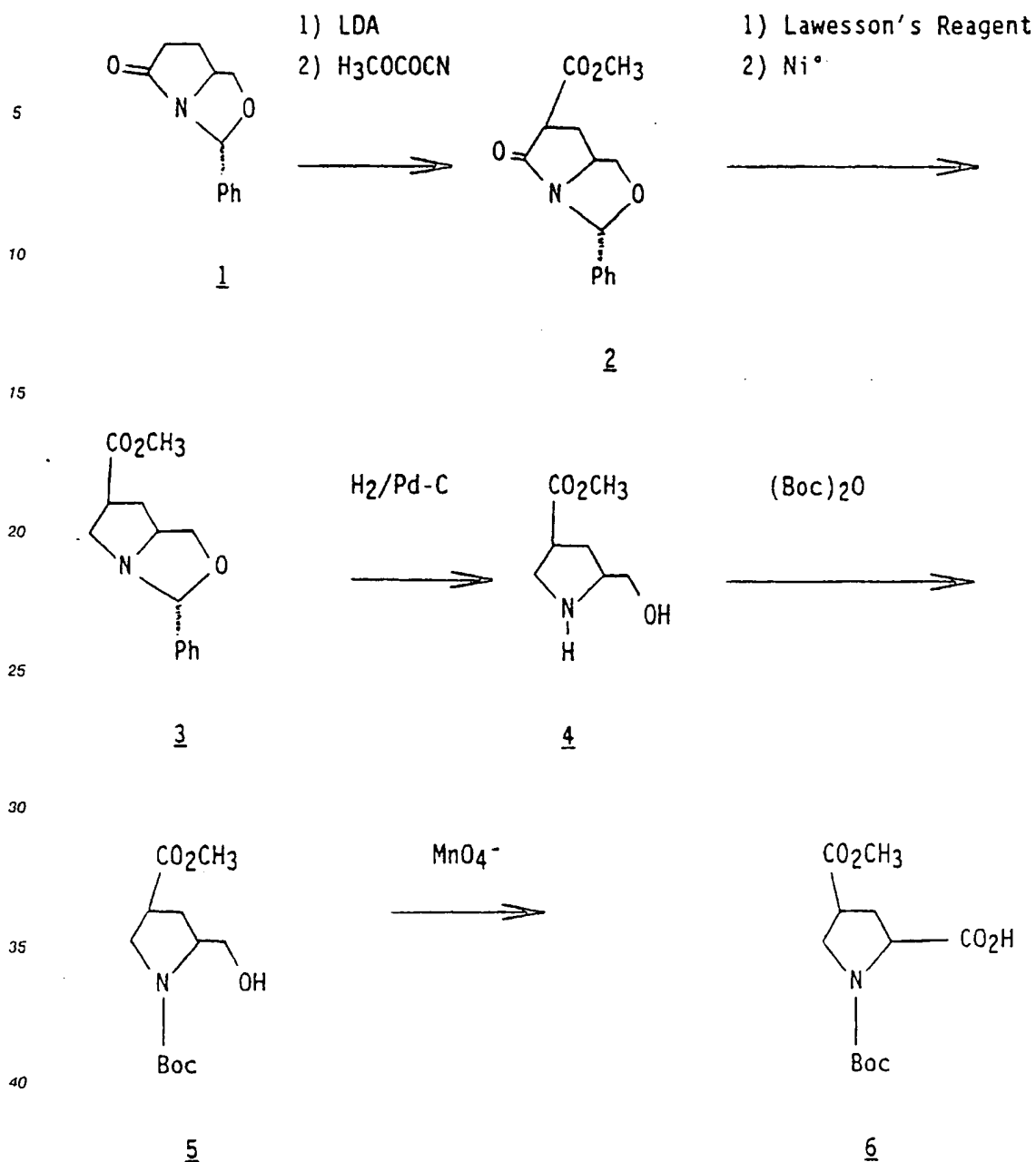
Preparation of H-Arg-Pro-Cpr-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Cpr³, DPhe⁷-bradykinin) and the preparation of the amino acid Cpr

Prior to the automated synthesis of the nonapeptide, the precursor for the unnatural amino acid residue
40 4-carboxyprolyl (Cpr) is prepared using the following procedure.

45

50

55



Methoxycarbonyl derivative (2):

The O,N-acetal (1) is added to a solution of lithium diisopropylamide in THF (prepared in the usual way from diisopropylamine and n-butyllithium) at -78 °C with vigorous stirring. After stirring another 30 minutes at -78 °C, a slight molar excess of methyl cyanoformate is added, and the temperature is raised to -15 °C with stirring for another hour. Quenching and purification give Compound 2.

(¹Thottathil, J.K., J.L. Moniot, R.H. Mueller, M.K.Y. Wong, T.P. Kissick, J. Org. Chem., Vol. 51, (1986) p. 3140.)

Protected prolinol derivative (3):

Compound 2 is treated with Lawesson's Reagent in warm THF to give the corresponding thiolactam. This is treated with Raney nickel in refluxing ethanol until reduction is complete. Purification gives Compound 3.

5

4-Methoxycarbonyl-L-prolinol (4):

Compound 3 is subjected to hydrogenolysis in methanol over 10% Pd on carbon at 45 psi until the starting material is consumed. Purification gives Compound 4.

10

N-Boc-4-methoxycarbonyl-L-prolinol (5):

Compound 4 is treated with (Boc)₂O in methanol with triethylamine at room temperature overnight. Purification gives compound 5.

15

N-Boc-4-Methoxycarbonyl-L-proline (6):

Compound 5 is oxidized with mildly alkylne permanganate in aqueous acetone. Purification gives Compound 6.

20

Compound 6 is incorporated into position 3 of the peptide sequence indicated in this example above using the same methods of peptide synthesis described in Example 1. The methyl ester of Compound 6 is cleaved by the final hydrogen fluoride cleavage of the peptide from the resin resulting in the amino acid residue Cpr in position 3 as shown. The structure of the nonapeptide is confirmed by mass spectral analysis, M + H = 1154, and the amino acid sequencing as described in Example 1.

25

Example 8

30

Preparation of H-Arg-Pro-Cmp-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Cmp³, DPhe⁷-bradykinin) and the preparation of the amino acid Cmp)

35

Prior to automated synthesis of the monapeptide, the precursor for the unnatural amino acid residue 4-carbomoxylprolyl (Cmp) is prepared as follows.

40

N-Boc-4-carbamoyl-L-proline (Boc-Cmp) (7):

Compound 6 from Example 7 is treated with methanol saturated with ammonia. Purification gives Compound 7.

45

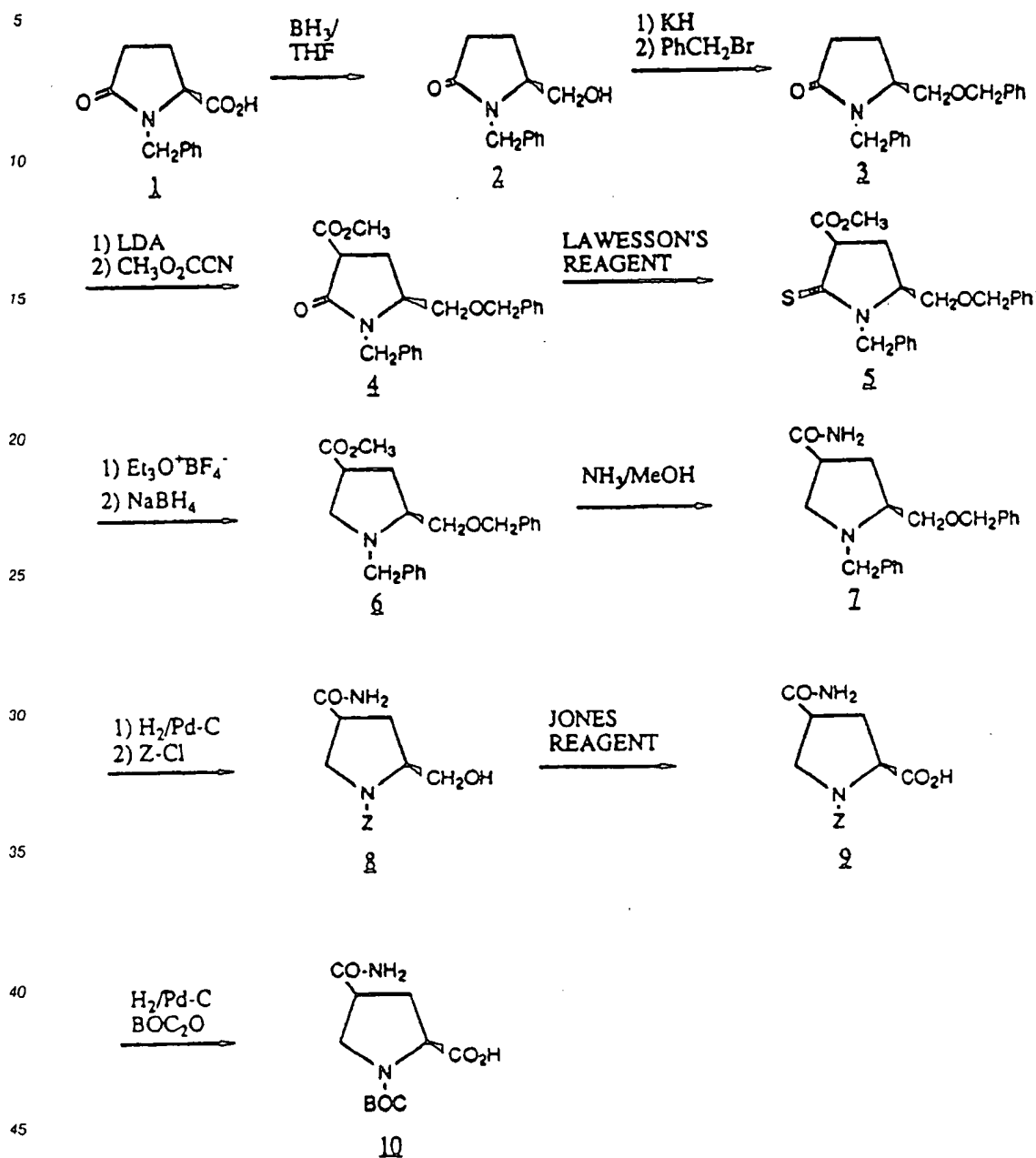
Compound 7 is incorporated into position 3 of the nonapeptide sequence indicated in this example above using the same methods of peptide synthesis described in Example 1. The structure of the nonapeptide is confirmed by mass spectral analysis, M + H = 1153, and the amino acid sequencing as described in Example 1.

50

Example 9

Preparation of H-Arg-Pro-Cmp-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Cmp³, DPhe⁷-bradykinin) and the preparation of the amino acid Cmp)

Prior to the automated synthesis of the nonapeptide, the precursor for the unnatural amino acid residue 4 carboxamidoproline (Cmp) is prepared using the following procedure.



⁵⁰ N-Benzyl pyroglutaminol (2):

A solution of borane in THF (1.4 equivalents) is added dropwise to a stirred solution of N-benzyl-L-pyroglutamic acid (1; See Petersen, J.S., G. Fels & H. Rapoport, Journal of the American Chemical Society, Vol. 106, (1984), p. 4539) in THF at 0°C, under argon. The solution is then stirred at room temperature overnight. The reaction is quenched by adding saturated aqueous K₂CO₃, then extracted with ethyl acetate. The extracts are dried and concentrated, and the residue is purified by chromatography (silica gel; CHCl₃/MeOH - 97/3). The product is crystallized from EtOAc/Hexane and has mp 83-84°C.

N-Benzyl pyroglutaminol benzyl ether (3):

A solution of Compound 2 in THF is added to a slight excess of potassium hydride suspended in THF at 0° C, under argon. After stirring for 1 hour, benzyl bromide is added, and the solution is warmed to room temperature and stirred another hour. The reaction is quenched with aqueous NaHCO₃, and extracted with ether. The extracts are dried and concentrated, and the residue is purified by chromatography (silica gel; CH₂Cl₂/EtOAc - 90/10). Compound 3 is obtained as an oil.

10 Methoxycarbonyl derivative (4):

Compound 3 is dissolved in THF and added to a solution of lithium diisopropylamine (2 equivalents) in THF (prepared in the usual way from diisopropylamine and n-butyllithium) at -78° C. The solution is warmed to 30° C over one hour, then cooled again to -78° C when methyl cyanoformate (1.05 equivalents) is added. After 10 minutes the solution is poured into 5% aqueous citric acid and extracted with EtOAc. The extracts are dried and concentrated and the residue is purified by chromatography to give Compound 4 as an oil.

20 Thiolactam (5):

Compound 4 is treated with Lawesson's Reagent in toluene at 60° C for 5 hours. Chromatography gives Compound 5 as an oil.

25 Protected prolinol derivative (6):

Compound 5 is treated with triethyloxonium tetrafluoroborate (1.1 equivalents) in CH₂Cl₂ for 1 hour at room temperature. The solvent is evaporated and replaced with methanol. Sodium borohydride (2 equivalents) is added, and the mixture is stirred for 1 hour. Purification provides Compound 6.

30 Carboxamide derivative (7):

Compound 6 is dissolved in ammonia saturated methanol and stirred for 3 days. Evaporation gives pure Compound 7.

40 Carbobenzoxy derivative (8):

Compound 7 is dissolved in glacial acetic acid treated with 10% palladium on carbon, and shaken under hydrogen (45 psi) until both benzyl groups are removed. The catalyst is removed by filtration and the solvent is evaporated. The residue is dissolved in water containing NaOH (3 equivalents) and treated with benzylchloroformate. The mixture is stirred vigorously for 4 hours, then extracted with EtOAc. The extracts are dried and concentrated, and the residue is purified by chromatography to give Compound 8.

45 N-Z-4-carboxamidoproline (9):

Compound 8 is dissolved in acetone and added to a stirred solution of Jones reagent in acetone at 0° C. After stirring several hours, isopropyl alcohol is added and stirring is continued another 20 minutes. After removing the organic solvents, the residue is diluted with water and extracted with EtOAc. The extracts are dried and concentrated to give Compound 9.

55 N-Boc-4-carboxyamidoproline (10):

The Z-group of Compound 9 is removed by hydrogenation in methanol with palladium/carbon catalyst. The N-Boc group is then introduced by stirring the secondary amine with Boc₂O and triethylamine in

aqueous THF. Reverse phase chromatography is required to provide pure Compound 10.

Compound 10 is incorporated into position 3 of the peptide sequence indicated above using the same methods of peptide synthesis described in Example 1. Hydrogen fluoride cleavage and purification as described above gives the nonapeptide. The structure of the nonapeptide is confirmed by mass spectral analysis, $M + H = 1153$ and the amino acid sequencing as described in Example 1.

Example 10

10

Preparation of H-Arg-Pro-Amp-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Amp³, DPhe⁷-bradykinin) and the preparation of α -N-Boc-N-Z-4-aminoproline

15

Prior to the automated synthesis of the nonapeptide, the precursor for the unnatural amino acid residue 4-aminoproline (Amp) is prepared using the following procedure.

α -N-Boc-N-Z-4-aminoproline:

N-Boc-4-aminoproline (see Example 6) is dissolved in a 1:1 mixture of dioxane and 2.5N NaOH. After chilling on ice, benzylchloroformate is added, and the mixture is stirred 30 minutes on ice, then 1 hour at room temperature. The mixture is diluted with water, washed with ether and then acidified to pH 2. Extraction with EtOAc and removal of solvent gives α -N-Boc-N-Z-4-aminoproline as a white solid.

The α -N-Boc-N-Z-4-aminoproline is incorporated into position 3 of the peptide sequence indicated above using the same methods of peptide synthesis described in Example 1. The Z-group is removed by the final hydrogen fluoride cleavage of the peptide from the resin resulting in the amino acid residue Amp in position 3 as shown. The structure of the nonapeptide is confirmed by mass spectral analysis, $M + H = 1125$, and amino acid sequencing as described in Example 1.

Example 11

35

Preparation of H-Arg-Pro-Aor-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Aor³, DPhe⁷-bradykinin)

Prior to the automated synthesis of the nonapeptide, the precursor for the amino acid residue Aor is prepared using the following procedure:

N(5)-Acetyl-N(2)-Boc-Ornithine (N-Boc-Aor):

45

To a solution of N(2)-Boc-ornithine (Bachem, Inc.) (2 mmol) and sodium carbonate (4 mmol) in water (10 mL) is added acetic anhydride (2.5 mmol). This mixture is stirred at room temperature for 3 hours, then acidified with solid citric acid and extracted several times with chloroform. Drying and removal of volatiles gives N-Boc-Aor.

The peptide is made using the same methods described in Example 1 with the replacement of N-Boc-Aor in the coupling reaction for position 3 of the peptide sequence. Cleavage and purification are accomplished as in Example 1 with the exception that the gradient for reverse phase chromatography is 20-35% solvent B (See Example 1). The purified peptide is analyzed by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1169$, and amino acid sequencing as described in Example 1.

Example 12

Preparation of H-DArg-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Arg-OH (DArg⁰, Asn³, DPhe⁷-bradykinin)

5

The peptide is made using the same methods described in Example 1 except that an additional residue of Boc-DArg(Tos) is added at the N-terminal end of the peptide following the Boc removal sequence described in Example 1. The final peptide is analyzed by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1283$, and amino acid sequencing as described in Example 1.

10

Example 13

15

Preparation of H-DArg-Arg-Pro-Aap-Gly-Phe-Ser-DPhe-Phe-Arg-OH (DArg⁰, Aap³, DPhe⁷-bradykinin)

20

The peptide is made using the same methods described in Example 1. The sequence is identical to that in Example 6 except that an additional residue of DArg is added to the N-terminus by coupling Boc-DArg(Tos) to the N-terminal end of the peptide following the Boc removal sequence described in Example 1. The final peptide is analyzed by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1323$, and amino acid sequencing as described in Example 1.

25

Example 14

30

Preparation of H-DArg-Arg-Pro-Cmp-Gly-Phe-Ser-DPhe-Phe-Arg-OH (DArg⁰, Cmp³, DPhe⁷-bradykinin)

The peptide is made using the same methods described in Example 1. The sequence is identical to that in Example 7 except that an additional residue of DArg is added to the N-terminus by coupling Boc-DArg(Tos) to the N-terminal end of the peptide following the Boc removal sequence described in Example 1. The final peptide is analyzed by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1309$, and amino acid sequencing as described in Example 1.

35

Example 15

40

Preparation of H-DArg-Arg-Pro-Nha-Gly-Phe-Ser-DPhe-Phe-Arg-OH (DArg⁰, Nha³, DPhe⁷-bradykinin)

45

The peptide is made using the same methods described in Example 1. N(4)-Hydroxyasparagine (Nha) is incorporated into the peptide by using N-Boc-aspartic acid β -methyl ester in the coupling reaction at position 3. DArg is incorporated into position 0 using Boc-DArg(Tos). The hydrogen fluoride cleavage leaves the methyl ester intact. After cleavage, the peptide is treated with hydroxylamine in aqueous solution to form the Nha residue. Purification is then accomplished as described in Example 1. The final peptide is analyzed by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1299$, and amino acid sequencing as described in Example 1.

50

Example 16

55

Preparation of H-Lys-Lys-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH (Lys-Lys⁰, Asn³, DPhe⁷-bradykinin)

The peptide is made using the same methods described in Example 1 and using the same sequence except after the Arg¹ is coupled the Boc removal sequence is again employed. Then N(2)-Boc-N(6)-CBZ-lysine is coupled, the peptide is cleaved as described in Example 1 and HPLC purification gives the peptide. The final peptide is analyzed by mass spectral analysis as described in Example 1 giving the correct molecular ion, $M + H = 1383$, and amino acid sequencing as described in Example 1.

METHODS FOR DETERMINING BRADYKININ ANTAGONIST ACTIVITY

The following test methods describe procedures used to measure the bradykinin antagonist activity of the peptides of the present invention. Different assays are considered predictive of the activity of the peptides on different tissues. A peptide which is an antagonist in one assay may be an agonist in other assays. Preferred bradykinin antagonist peptides are antagonists in all the predictive assays.

Test Method 1

Guinea Pig Ileum Assay

The bradykinin antagonists are assayed on guinea pig ileum for inhibition of the muscle-contracting activity of bradykinin, according to the commonly accepted assay method for bradykinin and related kinins as described in Stewart, J.M., "Chemistry and Biological Activity of Peptides Related to Bradykinin", Chapter V of Bradykinin, Kallidin and Kallikrein, (Handbook of Expt. Pharmacol.), E. G. Erdoes (ed.), Vol 25, Springer Verlag (pub.), 1970, pp. 242-243. The inhibition potencies are determined on isolated guinea pig ileum, according to the commonly accepted manner described for antagonists of biologically active compounds in Schild, H.O., "pA, a New Scale for the Measurement of Drug Antagonism", Br. J. Pharmacol., Vol. 2, No. 3 (1947), pp. 189-206). In the assay, a dose-response curve is determined for the reference substance bradykinin. The dose of bradykinin which produces a half maximal contraction of tissue is the ED₅₀ dose. An amount of bradykinin equivalent to twice the ED₅₀ dose is administered to the tissue 30 seconds after the start of incubation of the tissue with a dose of antagonist. Doses of antagonist are increased in this protocol until pre-incubation with a dose of antagonist reduces the contraction in response to the double ED₅₀ dose of bradykinin to response of a single ED₅₀ dose of bradykinin. The pA₂ value represents the negative logarithm of the molar concentration of antagonist which, in its presence, requires twice the amount of agonist to elicit the same response as when the agonist is given by itself. One unit of pA₂ value represents an order of magnitude change in potency. For comparison, the negative log of the dose of bradykinin, the dose which causes half maximal contraction of the tissues, is commonly known as the pD₂ value. The pD₂ value for bradykinin is 7.4 on the guinea pig ileum.

Test Method 2

Plasma Extravasation in the Guinea Pig Trachea

Male Hartley guinea pigs weighing 450-700 g are anesthetized with sodium pentobarbital. The right jugular vein is catheterized with a silastic tubing filled with 0.9% saline. The right carotid artery is cannulated. Evans blue dye is administered slowly via the jugular vein catheter. Five minutes later, bradykinin (0.4 ml/kg) and a test analogue (0.4 ml/kg) are co-administered intraarterially. Following an additional five minute period, the vasculature is perfused via the jugular vein with 0.9% saline. A 1 cm

section of tracheal tissue directly above the bifurcation is removed and weighed. Tissue is placed in tubes containing formamide. The tubes are placed in 50°C water-bath for 24 hours. Samples of the solvent from the tubes are transferred to cuvettes and measurements are taken on a fluorescence spectrophotometer. Calculations are made to determine dye content in the tissue. Data is expressed as ng dye/mg tissue.

5

Test Method 3

10

Mouse Abdominal Constriction Test

Compounds of the present invention are tested for analgesic activity using the mouse abdominal
15 constriction assay. This assay is described in Hendershot, L.C., & J. Forsaith, "Antagonism of the Frequency of Phenylquinone-induced Writhing in the Mouse by Weak Analgesics and Nonanalgesics", J. Pharmacol., Vol. 125, (1959), pp. 237-240, and in Methods in Narcotics Research, S. Ehrenpreis and A. Neidle, (eds.), Marcel Dekker, Inc., New York (pub.), 1975, pp. 64-65, the disclosures of both these references being incorporated herein in their entirety by reference.

20 Male CF-1 mice (Charles River Breeding Laboratories, Inc.), weighing approximately 20 grams and food fasted overnight, are used in these assays. Test compounds are prepared for intraperitoneal (i.p.) injection, with compounds of the present invention being prepared in distilled deionized water so that the appropriate dose is given in 0.2 mls to a 20 gram mouse.

25 An i.p. injection of 0.02% phenylquinone (2.5 mg/kg) is given at a concentration of 0.25 ml/20 grams body weight. Five minutes later the mice receive an i.p. injection of test compound. Five minutes after the i.p. injection of test compound, the number of full body writhes are counted for the succeeding ten minutes. Percent analgesia in this assay is calculated as follows:

$$30 \quad \frac{\text{Control number of writhes} - \text{Dosed number of writhes}}{\text{Control number of writhes}} \times 100$$

35 The mouse abdominal constriction assay was used to measure the antagonist activity of peptides of the present invention and related peptides as shown in Table 2:

Table 2

Mouse Abdominal Constriction Assay Results			
Compound No.	Peptide	Dose (mg/kg, i.p.)	Change from Control (%)
1	H-Arg-Pro-Hyp-Gly-Phe-Ser-DPhe-Phe-Arg-OH	25	-31
2	H-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH	50	57
3	H-Arg-Pro-Asn-DPro-Phe-Ser-DPhe-Phe-Arg-OH	50	72,61
4	H-Arg-Pro-Aap-Gly-Phe-Ser-DPhe-Phe-Arg-OH	50	46

50 Compound 1 is a peptide disclosed in U.S. Patent No. 4,693,993. The negative result in this assay indicates that this peptide is a bradykinin agonist according to this assay. Compounds 2, 3 and 4 are peptides of the present invention. These compounds all show bradykinin antagonist activity in the assay.

55

Test Method 4

Cell Culture Method5 Background.

In many different tissues and isolated cell populations, bradykinin induces a receptor mediated rise in the concentration of cytosolic free calcium ($[Ca^{2+}]_i$). In cultured NG108-15 neuronal cells (University of California at San Francisco, Cell Culture Facility, San Francisco, California), bradykinin increases $[Ca^{2+}]_i$ in a
 10 dose dependent manner from a basal concentration of about 260 nM to approximately 1000 nM at a maximally effective concentration of bradykinin (1×10^{-6} M). The half maximally effective concentration (EC_{50}) of bradykinin is 3×10^{-9} M.

Bradykinin receptor antagonist potency is evaluated by determining the concentration of antagonist required to inhibit by 50% (IC_{50}) the increase in $[Ca^{2+}]_i$ induced by 1×10^{-9} M bradykinin (Bachem,
 15 Torrance, CA).

NG108-15 Cell Culture.

20 NG108-15 cells are cultured in 75 cm³ Corning flasks (Fisher Scientific, Cincinnati, OH) in Dulbecco's modified minimal essential medium (high glucose; Gibco, Grand Island, NY) containing 5% fetal bovine serum (Hyclone, Logan, UT) and HAT (hypoxanthine-aminopterin-thymidine, Sigma, St. Louis, MO).

25 Measurement of Cytosolic Free Calcium Concentration in Cultured NG108-15 Cells

The concentration of cytosolic free Ca^{2+} ($[Ca^{2+}]_i$) is measured in suspensions of cultured NG108-15 cells loaded with the Ca^{2+} -sensitive fluorescent indicator fura-2 essentially as described by Osugi et al. (Journal Pharmacol. Exp. Ther., Vol. 240, (1987), pp. 617-622). Flasks of confluent cells are incubated for 60
 30 minutes at 37 °C in 20 ml of HEPES-buffered saline [140 mM NaCl, 5.4 mM KCl, 0.8 mM $MgSO_4$, 1.0 mM $CaCl_2$, 20 mM HEPES (N-2-hydroxyethylpiperazine-N'-2-ethanesulfonic acid; Behring Diagnostics, LaJolla, CA), 10 mM glucose, 0.25% bovine serum albumin (Cat. #A-4503; Sigma Chemical Co., St. Louis, MO)] containing fura-2 acetoxymethyl ester (Molecular Probes, Inc., Eugene, OR) at a final concentration of 2 μ M. The flasks are then washed twice with 30 ml of HEPES-buffered saline and the cells detached from the
 35 flasks in 15 ml of HEPES-buffered saline. The cell suspension is centrifuged at 25 °C for 5 minutes at 350 x g and the cell pellet resuspended to a final volume of approximately 10 ml of HEPES-buffered saline/flask. The cell suspension is maintained at room temperature prior to use.

The fluorescence of the fura-2 loaded NG108-15 cells is measured at an excitation wavelength of 340 nM (10 nM slit) and an emission wavelength of 500 nM (10 nM slit) in a Perkin-Elmer LS-5 fluorescence
 40 spectrophotometer fitted with a magnetic stirrer and a thermostatted cell holder. An aliquot of the cell suspension (1.37 ml) is placed in a cylindrical glass cuvette (internal diameter of 8 mm) containing a teflon coated stir bar and warmed to 37 °C in a water bath. After 5 minutes, the cuvette is placed in the cell holder of the fluorometer and stirring is initiated just prior to the addition of a test sample. Routinely, a test sample (0.014 ml of a 100-fold concentration solution) is added to the cuvette 30 seconds prior to the addition of 1×10^{-9} M bradykinin. Fluorescence is recorded continuously during the addition of the test sample and
 45 bradykinin.

Following completion of the fluorescence measurements, the cells are lysed with Triton X-100 at a final concentration of 10% (v/v) and the fluorescence of the Ca^{2+} saturated fura-2 (F_{max} ; fluorescence in the presence of 1 mM Ca^{2+}) and the Mn^{2+} -quenched fura-2 (autofluorescence (AF); fluorescence in the
 50 presence of 2 mM Mn^{2+}) determined. The $[Ca^{2+}]_i$ before and after the addition of the test sample and/or bradykinin is determined from the observed fluorescence values (F_{obs}) using these values of F_{max} and AF and the equations:

$$F_{min} = AF + (F_{max} - AF)/3.7$$

$$[Ca^{2+}]_i = K_D(F_{obs} - F_{min}) / (F_{max} - F_{obs})$$

55 where K_D is the dissociation constant for binding of Ca^{2+} to the dye (224 nM; Grynkiewicz et al., J. Biol. Chem., Vol. 260, (1985), pp. 3440-3450).

To quantitate the extent of inhibition by an antagonist, the peak increase in $[Ca^{2+}]_i$ from basal $[Ca^{2+}]_i$ (i.e., $[Ca^{2+}]_i$ just prior to the addition of bradykinin) is determined in the presence and absence of the

antagonist. The % inhibition is calculated as:

$$\frac{([Ca^{2+}] \text{ increase in presence of test sample})}{([Ca^{2+}] \text{ increase in absence of test sample})} \times 100$$

Agonistic activity is detected as an increase in fluorescence immediately following the addition of the test sample.

The cell culture method was used to measure the antagonist activity of peptides of the present invention and related peptides as shown in Table 3:

Table 3

Cell Culture Method Results		
Compound No.	Peptide	IC ₅₀
1	H-Arg-Pro-Hyp-Gly-Phe-Ser-DPhe-Phe-Arg-OH	Partial Agonist*
2	H-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH	5x10 ⁻⁷ M
5	H-DArg-Arg-Pro-Hyp-Gly-Phe-Ser-DPhe-Phe-Arg-OH	1x10 ⁻⁸ M (10% agonist*
6	H-DArg-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH	2.2x10 ⁻⁸ M

* Compounds 1 and 5 are peptides disclosed in U.S. Patent No. 4,693,993. Compound 1 exhibits high agonist (bradykinin-like) activity such that an antagonist potency could not be calculated from this assay. Compound 5 also exhibits small amount of agonist activity.

Compounds 2 and 6 are peptides of the present invention and both exhibit only antagonist (no agonist) activity in this assay.

Another aspect of the present invention includes methods of producing anti-inflammatory activity and/or analgesia or treating rhinitis, rhinorrhea and/or congestion in humans or lower animals by administering, to a human or lower animal in need of such present invention. This safe and effective amount can be given in a single dose or multiple doses repeatedly over the course of the treatment. Potential methods of administering the peptides of the present invention include per oral, parenteral (e.g., intravenous, intramuscular, intraperitoneal, subcutaneous), topical (epithelial, gingival, dermal or transdermal), intranasal, inhalation, sublingual, intravaginal, intrarectal, intrabuccal, or other internal administration means. Preferred methods of administering the peptides of this invention include dermal, transdermal, intranasal, and inhalation.

The phrase "safe and effective amount", as used herein, means an amount of a peptide or pharmaceutical composition of the present invention high enough to significantly positively modify the condition to be treated, but low enough to avoid serious side effects (at a reasonable benefit/risk ratio), within the scope of sound medical judgement. The safe and effective amount of the peptide or composition will vary with the particular condition being treated, the age and physical condition of the patient being treated, the severity of the condition, the duration of the treatment, the nature of concurrent therapy, the specific peptide or composition employed, the particular pharmaceutically-acceptable carrier utilized, and like factors within the knowledge and expertise of the attending physician.

The bradykinin antagonist peptides of this invention are useful for the treatment of one or more of the traumatic, inflammatory, pathological or normal conditions which are known to be mediated by bradykinin or exacerbated by an overproduction of bradykinin. These conditions involve pain, inflammation, congestion and/or smooth muscle contractions. Such conditions include wounds, burns, rashes, snake bites, insect bites and stings, angina, arthritis, asthma, allergies, rhinitis, rhinorrhea, sore throat pain, congestion, shock, inflammatory bowel disease, low blood pressure, systemic treatment of pain and inflammation, and/or low sperm motility which produces male infertility.

Therapeutic applications of the bradykinin antagonist peptides of the present invention include not only treatment for the production of bradykinin or related kinins by the animal but also the injection of bradykinin related peptides into an animal as a result of bites and stings. Topical application alone or in combination with subcutaneous utilization of the bradykinin antagonists of the invention can be employed to treat the effects of bradykinin-related peptides causing pain, inflammation, swelling and/or smooth muscle contraction.

It is estimated that the dosage range for topical application in such conditions as the pain and inflammation of wound, burns, rashes and sore throat would be from about 0.1 ml to about 500 mg per dose, preferably from about 1 mg to about 10 mg per dose. For a nasal or inhalation spray formulation suitable for treating rhinitis, allergies and asthma, a suitable dosage range would be from about 0.01 mg to about 100 mg per dose, preferably from about 0.5 mg to about 5 mg per dose. For an intravenous formulation suitable for the treatment of systemic inflammation, shock, arthritis, allergies, asthma and for increasing sperm motility, a suitable dosage range would be from about 0.1 mg/kg to about 500 mg/kg per dose, preferably from about 5 mg/kg to about 50 mg/kg per dose. For a per oral formulation for the treatment of inflammatory bowel disease or general pain and inflammation, a suitable dosage range would be from about 1 mg/kg to about 1000 mg/kg per dose, preferably from about 50 mg/kg to about 500 mg/kg per dose.

Another aspect of the present invention involves pharmaceutical compositions comprising a peptide of the present invention and a pharmaceutically-acceptable carrier. The peptides of the present invention typically comprise from about 0.1% to about 10% by weight of the pharmaceutical compositions of the present invention, preferably from about 0.5% to about 5%, and most preferably from about 1% to about 3%.

The term "pharmaceutically-acceptable carrier", as used herein, means one or more compatible solid or liquid filler diluents or encapsulating substances which are suitable for administration to a human or lower animal. The term "compatible", as used herein, means that the components of the pharmaceutical compositions are capable of being commingled with the compound of the present invention, and with each other, in a manner such that there is no interaction which would substantially reduce the pharmaceutical efficacy of the pharmaceutical composition under ordinary use situations. Pharmaceutically-acceptable carriers must, of course, be of sufficiently high purity and sufficiently low toxicity to render them suitable for administration to the human or lower animal being treated.

Some examples of substances which can serve as pharmaceutically-acceptable carriers are sugars such as lactose, glucose and sucrose; starches such as corn starch and potato starch; cellulose and its derivatives, such as sodium carboxymethylcellulose, ethylcellulose, cellulose acetate; powdered tragacanth; malt; gelatin; talc; stearic acid; magnesium stearate; calcium sulfate; vegetable oils such as peanut oil, cottonseed oil, sesame oil, olive oil, corn oil and oil of theobroma; polyols such as propylene glycol, glycerine, sorbitol, mannitol, and polyethylene glycol; sugar; alginic acid; pyrogen-free water; isotonic saline; phosphate buffer solutions; cocoa butter (suppository base); emulsifiers, such as the Tweens®; as well as other non-toxic compatible substances used in the pharmaceutical formulations. Wetting agents and lubricants such as sodium lauryl sulfate, as well as coloring agents, flavoring agents, excipients, tableting agents, stabilizers, antioxidants, and preservatives, can also be present.

While particular embodiments of the present invention have been described, it will be obvious to those skilled in the art that various changes and modifications to the peptides and compositions disclosed herein can be made without departing from the spirit and scope of the invention. It is intended to cover, in the appended claims, all such modifications that are within the scope of this invention.

Claims

1. A bradykinin antagonist peptide characterized in that it has the structure:
H-L-Q-T-U-V-W-X-Y-Z-Arg-OH

wherein

- (a) L is selected from nil, a single basic amino acid residue of the D- or L-configuration, a peptide containing from 2 to about 4 amino acid residues of the D-or L-configuration, at least one of which is a basic amino acid residue, and an N-terminal enzyme protecting group from the group consisting of acyl-type protecting groups, aromatic urethane-type protecting groups, and alkyl-type protecting groups;
- (b) Q is selected from Arg, Lys and Orn;
- (c) T and U are each independently selected from D- and L-cyclic amino acid residues, D- and L-aliphatic amino acid residues, L-aromatic amino acid residues, L-acidic amino acid residues, L-amide amino acid residues and L-hydroxamate amino acid residues; wherein at least one of T and U is selected from L-acidic amino acid residues, L-amide amino acid residues and L-hydroxamate amino acid residues;
- (d) V is selected from aliphatic, cyclic and aromatic amino acid residues of the D- or L-configuration;
- (e) W is selected from aliphatic and aromatic amino acid residues of the L-configuration;
- (f) X is selected from aromatic and aliphatic amino acid residues of the D- or L-configuration;

(g) Y is selected from aromatic amino acid residues of the D-configuration;
 (h) Z is selected from aromatic and aliphatic amino acid residues of the L-configuration;
 and the pharmaceutically-acceptable salts thereof.

2. The peptide of Claim 1 characterized in that

- 5 (a) L is selected from nil, Arg, DArg, Lys, DLys, Lys-Lys, Met-Lys, Gly-Arg-Met-Lys, DLys-Lys, Orn, Har, DHar, 5-guanidinopentanoyl, 6-guanidinohexanoyl, 6-aminohexanoyl and 5-aminopentanoyl;
 (b) Q is selected from Arg, Lys, and Orn;
 (c) T and U are each independently selected from Pro, DPro, Hyp, Azt, Thz, Inp, Epr, Asp, Glu, Asn, Gln, Cpr, Cmp, Aap, Aor, Nha and Nhg; wherein at least one of T and U is selected from Asp, Glu, Asn,
 10 Gln, Cpr, Cmp, Aap, Aor, Nha and Nhg;
 (d) V is selected from Gly, Ala, DPro, DHyp and Sar;
 (e) W is selected from Phe, Thi, Nap, Pyr, Nph, Cph, Mty and Tyr;
 (f) X is selected from Ser, Thr, Ala, Gly, DPhe, DThi, DNap, DPyr and DCph;
 (g) Y is selected from DPhe, DTyr, DThi, DNap, DPyr, DNph, DCph, DMty, DTrp, DHis, DHph and
 15 DPgl;
 (h) Z is selected from Phe, Tyr, Thi, Pyr, Nap, Mty, Nph and Cph.

3. The peptide of Claim 2 wherein

- (a) L is selected from DArg, Lys-Lys, and nil; preferably L is DArg;
 20 (b) Q is Arg;
 (c) T and U are each independently selected from Pro, Hyp, Asp, Glu, Asn, Gln, Cpr, Cmp, Aap, Aor, Nha and Nhg; wherein at least one of T and U is selected from Asp, Glu, Asn, Gln, Cpr, Cmp, Aap, Aor, Nha and Nhg;
 (d) V is selected from DPro, DHyp, Gly and Ala; preferably V is Gly or DPro;
 25 (e) W is selected from Phe and Tyr; preferably W is Phe;
 (f) X is selected from Ser, Thr, Ala and Gly; preferably X is Ser;
 (g) Y is selected from DPhe, DTyr, DTrp and DHis; preferably Y is DPhe; and
 (h) Z is selected from Phe and Tyr; preferably Z is Phe.

30 4. The peptide of any of Claims 1-3 wherein one of T or U, preferably T, is selected from Pro and Hyp; and the other of T or U, preferably U, is selected from Asp, Glu, Asn, Gln, Cpr, Cmp, Aap, Aor, Nha and Nhg, preferably from Asp, Glu, Aap, Asn and Gln, most preferably from Aap, Asn and Gln.

5. The peptide of Claim 2 characterized in that it has a structure wherein:

- (a) Q is Arg;
 35 (b) T is Pro or Hyp;
 (c) U is Aap, Asn or Gln;
 (d) V is Gly or DPro;
 (e) W is Phe;
 (f) X is Ser;
 40 (g) Y is DPhe;
 (h) Z is Phe; and
 (i) L is DArg.

6. A peptide characterized in that it has a structure selected from:

- 45 H-Arg-Pro-Asn-DPro-Phe-Ser-DPhe-Phe-Arg-OH,
 H-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH,
 H-DArg-Arg-Pro-Asn-DPro-Phe-Ser-DPhe-Phe-Arg-OH,
 H-DArg-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH,
 H-Lys-Lys-Arg-Pro-Asn-DPro-Phe-Ser-DPhe-Phe-Arg-OH,
 50 H-Lys-Lys-Arg-Pro-Asn-Gly-Phe-Ser-DPhe-Phe-Arg-OH,
 H-Arg-Pro-Aap-DPro-Phe-Ser-DPhe-Phe-Arg-OH,
 H-Arg-Pro-Aap-Gly-Phe-Ser-DPhe-Phe-Arg-OH,
 H-DArg-Arg-Pro-Aap-DPro-Phe-Ser-DPhe-Phe-Arg-OH,
 H-DArg-Arg-Pro-Aap-Gly-Phe-Ser-DPhe-Phe-Arg-OH,
 55 H-Lys-Lys-Arg-Pro-Aap-DPro-Phe-Ser-DPhe-Phe-Arg-OH and
 H-Lys-Lys-Arg-Pro-Aap-Gly-Phe-Ser-DPhe-Phe-Arg-OH.

7. A pharmaceutical composition characterized in that it comprises the peptide of any of Claims 1-6 and a pharmaceutical carrier.

8. The use of a safe and effective amount of the peptide of any of Claims 1-6 for the manufacture of a medicament to treat pain, inflammation, congestion or smooth muscle contraction.

5

10

15

20

25

30

35

40

45

50

55

19



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) Publication number:

0 334 244 A3

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 89104872.0

(51) Int. Cl.⁵: **C07K 7/18, A61K 37/02,
//C07K99/18**

(22) Date of filing: 18.03.89

(30) Priority: 25.03.88 US 173391
01.03.89 US 315245

(43) Date of publication of application:
27.09.89 Bulletin 89/39

(94) Designated Contracting States:
AT BE CH DE FR GB IT LI LU NL SE

(98) Date of deferred publication of the search report:
29.05.91 Bulletin 91/22

(71) Applicant: **THE PROCTER & GAMBLE
COMPANY**
One Procter & Gamble Plaza
Cincinnati Ohio 45202(US)

(72) Inventor: **Gardner, Joseph Herman**
4060 Boomer Road
Cincinnati, OH 45247(US)
Inventor: **Correa, Paul Elliott**
6511 Wesselman Road
Cincinnati, OH 45248(US)
Inventor: **Berman, Elizabeth Faith**
11787 Rose Lane
Cincinnati, OH 45246(US)
Inventor: **Charest, Robert Paul**
3935 Dust Commander Drive
Hamilton, OH 45011(US)

(74) Representative: **Canonici, Jean-Jacques et al**
Procter & Gamble European Technical
Center N.V. Temseilaan 100
B-1853 Strombeek-Bever(BE)

(54) **Bradykinin Antagonist Peptides.**

(57) The present invention involves nona- or larger bradykinin antagonist peptides characterized by the structure: H-L-Q-T-U-V-W-X-Y-Z-Arg-OH where certain amino acid residues of the nonapeptide hormone bradykinin or other substituted analogs of bradykinin are altered as follows: either position 2, position 3, or both consist of a L-acidic, L-amide or L-hydroxamate amino acid residue; position 7 consists of a D-aromatic amino acid residue; and position 4 preferably consists of a L-aliphatic or a D-cyclic amino acid residue.

EP 0 334 244 A3



EUROPEAN SEARCH REPORT

EP 89 10 4872

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
X	WO-A-8 607 263 (STEWART) * Completely * - - - -	1-2,4,6-7	C 07 K 7/18 A 61 K 37/02 // C 07 K 99/18
P,X	WO-A-8 901 781 (STEWART) * Completely * - - - -	1-2,4,6-7	
P,X	WO-A-8 901 780 (STEWART) * Completely * - - - -	1-2,4,6-7	
X	PEPTIDES, STRUCTURES AND FUNCTION, PROCEEDINGS OF THE NINTH AMERICAN PEPTIDE SYMPOSIUM, edited by C.M. DEBER, pages 655-658, Pierce Chemical Co., Rockford, Illinois, US; R.J. VAVREK et al.: "Development and modification of competitive antagonists of bradykinin" * Complete article * - - - -	1-2,4,6-7	
A	US-A-3 234 200 (M. BODANSZKY) * Completely * - - - - -	1,7	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C 07 K A 61 K
The present search report has been drawn up for all claims			
Place of search		Date of completion of search	Examiner
The Hague		26 February 91	RAJIC M.
<div>CATEGORY OF CITED DOCUMENTS</div> <div><div>X: particularly relevant if taken alone Y: particularly relevant if combined with another document of the same category A: technological background O: non-written disclosure P: intermediate document T: theory or principle underlying the invention</div><div>E: earlier patent document, but published on, or after the filing date D: document cited in the application L: document cited for other reasons &: member of the same patent family, corresponding document</div></div>			